

## **BILLING CODE 4410-09-P**

## **DEPARTMENT OF JUSTICE**

**Drug Enforcement Administration** 

[Docket No. DEA-392]

Importer of Controlled Substances Application: Akorn, Inc.

**ACTION:** Notice of application.

DATES: Registered bulk manufacturers of the affected basic classes, and applicants therefore, may file written comments on or objections to the issuance of the proposed registration on or before [INSERT DATE 30 DAYS AFTER DATE OF PUBLICATION IN THE FEDERAL REGISTER]. Such persons may also file a written request for a hearing on the application on or before [INSERT DATE 30 DAYS AFTER DATE OF PUBLICATION IN THE FEDERAL REGISTER].

ADDRESSES: Written comments should be sent to: Drug Enforcement Administration, Attention: DEA Federal Register Representative/DPW, 8701 Morrissette Drive, Springfield, Virginia 22152. All requests for hearing must be sent to: Drug Enforcement Administration, Attn: Administrator, 8701 Morrissette Drive, Springfield, Virginia 22152. All requests for hearing should also be sent to: (1) Drug Enforcement Administration, Attn: Hearing Clerk/OALJ, 8701 Morrissette Drive, Springfield, Virginia 22152; and (2) Drug Enforcement Administration, Attn: DEA Federal Register Representative/DPW, 8701 Morrissette Drive, Springfield, Virginia 22152.

SUPPLEMENTARY INFORMATION: The Attorney General has delegated his

authority under the Controlled Substances Act to the Administrator of the Drug

Enforcement Administration (DEA), 28 CFR 0.100(b). Authority to exercise all

necessary functions with respect to the promulgation and implementation of 21 CFR part

1301, incident to the registration of manufacturers, distributors, dispensers, importers,

and exporters of controlled substances (other than final orders in connection with

suspension, denial, or revocation of registration) has been redelegated to the Assistant

Administrator of the DEA Diversion Control Division ("Assistant Administrator")

pursuant to section 7 of 28 CFR part 0, appendix to subpart R.

In accordance with 21 CFR 1301.34(a), this is notice that on May 2, 2018, Akorn Inc.,

1222 W. Grand Avenue, Decatur, Illinois 62522 applied to be registered as an importer of

the following basic class of controlled substance:

**Controlled Substance Drug Code Schedule** Remifentanil 9739  $\Pi$ 

The company plans to import the above listed controlled substance for research

purposes.

Dated: November 16, 2018.

John J. Martin,

Assistant Administrator.

[FR Doc. 2018-25873 Filed: 11/26/2018 8:45 am; Publication Date: 11/27/2018]

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